



IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Application No. : 10/625,626                      Art unit : 1625  
Applicant : Steven P. Adams et al.              Examiner : Janet L. Coppins  
Filing date : July 24, 2003  
Title : CELL ADHESION INHIBITORS

Mail Stop AF  
Customer Service Window  
Randolph Building  
401 Dulaney Street  
Alexandria, VA 22314

DO NOT  
ENTER  
JLC

**REPLY TO OFFICE ACTION MAILED JANUARY 13, 2005**

In response to the Office Action mailed January 13, 2005 ("Office Action"), Applicants submit the following remarks.

**REMARKS**

Applicants thank the Examiner for indicating that claims 1-7, 10 and 11 are allowable. Claims 12-15 stand rejected.

**Rejection under 35 U.S.C. § 112, first paragraph**

Claims 12-15 have been rejected by the Examiner under 35 U.S.C. § 112, first paragraph, for lack of enablement. Claim 12 is independent, and claims 13-15 depend from it.

The Examiner contends that claims 12-15 are not supported by a specific asserted utility or a well established utility that would enable one skilled in the art to know how to use the claimed invention. See the Office Action at page 2. Specifically, the Examiner contends that because "claim 12 is not directed to a method of treating a disease," the Applicants have failed to set forth a definable utility. See the Office Action at page 2.

Independent claim 12 is directed to a method of preventing, inhibiting or suppressing cell adhesion in a mammal in need thereof. To satisfy enablement under 35 U.S.C. § 112, "All that is necessary is that one skilled in the art be able to practice the *claimed* invention, given the level of knowledge and skill in the art." (emphasis added) MPEP 2164.08; *see e.g. In re Fisher*, 427 F.2d 833, 839, 166 USPQ 18, 24 (CCPA 1970). At the time of filing, one of ordinary skill in the art would have been apprised of the disclosed article, Lobb, R.R. and Hemler, M.E., "The Pathophysiologic Role of  $\alpha 4$  Integrins In Vivo," *J. Clin. Invest.*, 94, pp. 1722-29 (1994) ("Lobb"), which indicates that "rapidly accumulating in vivo data... suggest that  $\alpha 4$  integrin-